



Sensitive Inexpensive HPLC-UV Method for Estimation of Cyproterone Acetate in Pharmaceuticals

Shahid SHAH¹, Muhammad HANIF², Muhammad Ahmad SHAHZAD³, Malik SAADULLAH⁴, Khalid MAHMOOD⁵, Ayesha ASLAM⁶, Zubaida BI BI⁵, Nazar ABBAS⁷, Muhammad Masood AHMED⁸, Sajid Mehmood KHAN⁹, Ghulam ABBAS^{7*}, Omeira IQBAL⁷ & Mehran ASHFAQ⁷

¹ Department of Pharmacy Practice, ⁴ Department of Pharmaceutical Chemistry,

⁷ Department of Pharmaceutics, Faculty of Pharmaceutical Sciences,
Government College University Faisalabad, Pakistan.

² Department of Pharmaceutics, Faculty of Pharmacy, Bahauddin Zakariya University Multan, Pakistan

³ Department of Statistics, ⁵ Institute of Chemical Sciences,

⁸ Faculty of Pharmacy, Bahauddin Zakariya University Multan, Pakistan

⁶ Department of Neurology, King Edward Medical University Lahore, Pakistan

⁹ Faculty of Pharmacy and Alternative Medicine, The Islamia University Bahawalpur, Pakistan

SUMMARY. A simple, precise and inexpensive HPLC-UV method has been developed for the estimation of cyproterone acetate using high performance liquid chromatography (HPLC) on C₁₈ column with UV detection at 280 nm. The expected optimal assay condition comprised of acetonitrile and phosphate buffer pH 5.8 in ratio of 40:60 % v/v at a flow rate of 1.5 mL/min. Under this optimal state, detection of CA with good resolution and retention time less than 5 min were attained. The method was proved to be linear in the range of 200 to 600 ng/mL and 0.9996 regression values was obtained for CA. The percentage recovery of CA was ranged from 98.25 to 99.17 %. The precision and selectivity of the developed method was good and can be used for the estimation of CA in pharmaceuticals.

RESUMEN. Se ha desarrollado un método HPLC-UV simple, preciso y económico para la estimación del acetato de ciproterona mediante cromatografía líquida de alta resolución (HPLC) en columna C₁₈ con detección UV a 280 nm. La condición de ensayo óptima esperada comprendía acetonitrilo y tampón fosfato pH 5,8 en una proporción de 40:60% v/v a un caudal de 1,5 mL/min. En este estado óptimo, se logró la detección de CA con buena resolución y un tiempo de retención de menos de 5 min. Se demostró que el método es lineal en el rango de 200 a 600 ng/mL y se obtuvieron valores de regresión de 0.9996 para CA. El porcentaje de recuperación de CA varió de 98,25 a 99,17%. La precisión y selectividad del método desarrollado fue buena y se puede utilizar para la estimación de CA en productos farmacéuticos.

KEY WORDS: cyproterone acetate, HPLC, pharmaceuticals.

* Author to whom correspondence should be addressed. E-mail: ghulamabbas@gcuf.edu.pk